This listing of claims will replace all prior versions, and listing, of claims in the application.

Listing of Claims:

- 1. (Currently Amended) A pharmaceutical composition comprising metaxalone in a pharmaceutically acceptable solubility improved form and at least one pharmaceutically acceptable excipient, characterized in that the pharmaceutical composition has enhanced oral bioavailability a greater rate and extent of absorption as compared to the pharmaceutical composition of metaxalone described in New Drug Application No. 13-217 when they are orally administered to a patient on an empty stomach, wherein at least 99% of the metaxalone has a particle size not more than about 10µm in diameter.
 - 2. (Cancelled)
 - 3. (Cancelled)
 - 4. (Cancelled)
 - 5. (Cancelled)
 - 6. (Cancelled)
- 7. **(Original)** A pharmaceutical composition as claimed in claim 1, wherein the composition comprises a mixture of metaxalone and a solubilizing agent.
 - 8. (Cancelled)
 - 9. (Cancelled)
- 10. (Currently Amended) A pharmaceutical composition as claimed in claim 9 1, wherein the metaxalone has specific surface area per unit volume of more than 2.5m²/cm³.
- 11. **(Previously Presented)** A pharmaceutical composition as claimed in claim 10, wherein the metaxalone has specific surface area per unit volume of more than $3.0 \text{m}^2/\text{cm}^3$.
 - 12. (Cancelled)

Application No. 10/526,285 Amendment dated October 27, 2008 Response to Office Action dated March 20, 2008

13. (Cancelled)

- 14. (Currently Amended) A pharmaceutical composition as claimed in claim 1, wherein the metaxalone comprises the following particle size distribution characteristics: 99% undersize value of $10\mu m$, 90% undersize value of $5\mu m$, and 50% undersize value of $3\mu m$.
- 15. **(Previously Presented)** A pharmaceutical composition as claimed in claim 1, wherein the amount of metaxalone is in the range of 400mg to 1600mg.
- 16. (Previously Presented) A pharmaceutical composition as claimed in claim 1, wherein the pharmaceutically acceptable excipient comprises a wetting agent.
- 17. **(Previously Presented)** A pharmaceutical composition as claimed in claim 16, wherein the wetting agent comprises a surfactant.
- 18. **(Previously Presented)** A pharmaceutical composition as claimed in claim 17, wherein the surfactant comprises sodium lauryl sulfate.

19-22 (Cancelled)

- 23. (Previously Presented) A pharmaceutical composition as claimed in claim 1 wherein the pharmaceutical composition further comprises an analgesically effective amount of a non-steroidal anti-inflammatory drug, wherein said nonsteroidal anti-inflammatory drug comprises a propionic acid derivative, acetic acid derivative, fenamic acid derivative, biphenylcarboxylic acid derivative or an oxicam, or the pharmaceutically acceptable salts thereof.
 - 24. (Cancelled)
 - 25. (Cancelled)
 - 26. (Cancelled)
 - 27. (Cancelled)
 - 28. (Cancelled)

Application No. 10/526,285 Amendment dated October 27, 2008

Response to Office Action dated March 20, 2008

29. (Currently Amended) A method of relieving discomfort associated with

acute, painful musculo-skeletal conditions comprising orally administering to a patient a

pharmaceutical composition comprising metaxalone in a pharmaceutically acceptable solubility-

improved form and at least one pharmaceutically acceptable excipient, characterized in that the

pharmaceutical composition has enhanced oral bioavailability a greater rate and extent of

absorption as compared to the pharmaceutical composition of metaxalone described in New

Drug Application No. 13-217 when they are orally administered to a patient on an empty

stomach, wherein at least 99% of the metaxalone has a particle size not more than about 10 µm in

diameter.

30. (New) A method comprising orally administering to a patient a pharmaceutical

composition comprising metaxalone and at least one pharmaceutically acceptable excipient,

characterized in that the pharmaceutical composition has a greater rate and extent of absorption

as compared to the pharmaceutical composition of metaxalone described in New Drug

Application No. 13-217 when orally administered to a patient on an empty, wherein the

metaxalone has specific surface area per unit volume of more than about 2.5m²/cm³.

31. (New) A pharmaceutical composition comprising metaxalone and at least one

pharmaceutically acceptable excipient, characterized in that the pharmaceutical composition has

a greater rate and extent of absorption as compared to the pharmaceutical composition of

metaxalone described in New Drug Application No. 13-217 when orally administered to a

patient on an empty, wherein the metaxalone has specific surface area per unit volume of more

than about $2.5 \text{m}^2/\text{cm}^3$.

6